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Substitute for form 1449A/PTO INFORMATION DISCLOSURE STATEMENT BY APPLICANT (use as many sheets as necessary) Sheet 1 of 3	Application Number	10/690,115
	Filing Date	October 21, 2003
	First Named Inventor	Richard Apodaca
	Group Art Unit	1624
	Examiner Name	COLEMAN
	Attorney Docket Number	PRD 2033 NP

U.S. PATENT DOCUMENTS

Examiner Initials	Cite No. ¹	U.S. Patent Document		Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document mm-dd-yyyy	Pages, Columns, Lines, where relevant passages or relevant figures appear	
		Number	Kind Code ² (if known)				
BC		USPN 3,888,160		Tweit, Robert C.	05-27-1975	544	316
BC		USPN 3,714,179		Tweit, Robert C.	01-30-1973	548	315.7
BC		USPN 5,030,644		Baldwin et al.	07-09-1991	514	393
BC		USPN 5,217,986		Pomponi, S.A. et al.	06-08-1993	514	400
BC		USPN 5,352,707		Pompri, S.A. et al.	10-04-1994	514	651
BC		USPN 5,869,479		Kreutner, W.; Hey, J.A.	02-09-1999	514	217.05

FOREIGN PATENT DOCUMENTS

Examiner Initials	Cite No. ¹	Foreign Patent Document			Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document mm-dd-yyyy	Pages, Columns, Lines, where relevant passages or relevant figures appear	T ³
		Office ³	Number ⁴	Kind Code ⁵				
BC	✓	WO	99/42458		James Black Foundation Limited	08-28-1999		
BC	✓	EP	0878512	A1	Societe Civile Bioprojet	02-09-2000		
** BC	✓	JP	02308237	A2	Kato et al.	12-19-1990		
BC	✓	WO	02/076925	A2	Eli Lilly and Company	10-03-2002		
BC	✓	WO	03/050099	A1	Ortho-McNeil Pharmaceutical, Inc.	06-18-2003		
BC	✓	WO	02/024695	A2	Ortho-McNeil Pharmaceutical, Inc.	03-28-2002		
BC	✓	WO	02/012214	A2	Ortho-McNeil Pharmaceutical, Inc.	02-14-2002		
BC	✓	WO	02/012190	A2	Ortho-McNeil Pharmaceutical, Inc.	02-14-2002		
BC	✓	WO	03/064411	A1	Novo Nordisk	08-07-2003		
BC	✓	WO	03/031432	A1	Novo Nordisk	04-17-2003		
BC	✓	WO	03/024929	A1	Novo Nordisk	03-27-2003		
BC	✓	WO	03/004480	A2	Novo Nordisk	01-16-2003		
BC	✓	WO	03/024928	A2	Novo Nordisk	03/27/2003		

Examiner Signature	Brenda Coleman	Date Considered	May 30, 2006
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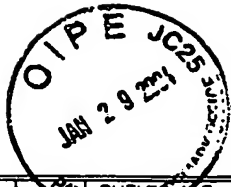
INFORMATION DISCLOSURE STATEMENT BY APPLICANT

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Sheet 2 of 3

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First Named Inventor	Richard Apodaca
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OTHER PRIOR ART - NON PATENT LITERATURE DOCUMENTS			
Examiner's Initials*	Cite No. ¹	Include name of the author (in CAPITOL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published	T ²
BC	✓	ALBENGRES, E. et al. Systemic Antifungal Agents. Drug Safety (Feb. 1998) 18(2):83-97	
BC	✓	ALI, S.M. et al. Design, Synthesis, and Structure-Activity Relationships of Acetylene-Based Histamine H3 Receptor Antagonists. J. Med. Chem. (1999) 42(5):903-909	
BC	✓	ARRANG, J.-M. et al. Auto-inhibition of Brain Histamine Release Mediated by a Novel Class (H3) of Histamine Receptor. Nature (April 1983) 302:832-837	
BC	✓	ASH, A.S.F.; SCHILD, H.O. Receptors Mediating Some Actions of Histamine. Br. J. Pharmac. Chemother. (1968) 27:427-439	
BC	✓	BACK, D.J.; TJIA, J.F. Inhibition of Tolbutamide Metabolism by Substituted Imidazole Drugs In Vivo: Evidence for a Structure-Activity Relationship. Br. J. Pharmacol. (1985) 85:121-126	
BC	✓	BARNES, J.C. et al. The Selective Histamine H3 Receptor Antagonist Thioperamide Improves Cognition and Enhances Hippocampal Acetylcholine Release In Vivo. Soc. Neurosci. Abstr. (1993) 19:1813	
BC	✓	Bioworld Today, March 2, 1999, page 3	
BC	✓	BLACK, J.W. et al. Definition and Antagonism of Histamine H2-Receptors. Nature (April 1972) 238:385-390	
BC	✓	DING, Y.-S. et al. Synthesis of High Specific Activity (+)- and (-)-8-[18F]Fluoronoropinephrine via the Nucleophilic Aromatic Substitution Reaction. J. Med. Chem. (1991) 34(2):767-771	
BC	✓	GANELLIN, C.R. et al. Synthesis of Potent Non-Imidazole Histamine H3-Receptor Antagonists. Arch. Pharm. Pharm. Med. Chem. (Weinheim, Ger.) (1998) 331:395-404	
BC	✓	GARBARG, M. et al. S-[2-(4-Imidazolyl)ethyl]isothiourea, a Highly Specific and Potent Histamine H3 Receptor Agonist. J. Pharmacol. Exp. Ther. (1992) 263(1):304-310	
BC	✓	Gilatech Inc. Press Release Nov. 5, 1998	
BC	✓	GONZALEZ, F. GARCIA, et al. Synthesis of 3-aryl(alkyl)-4-(D-arabino-tetrahydroxybutyl)imidazoline-2-thiones. Carbohydrate Research, 22(2), 438-40 (English) 1988	
BC	✓	ICHINOSE, M.; BARNES, P.J. Histamine H3-Receptors Modulate Nonadrenergic Noncholinergic Neural Bronchoconstriction in Guinea-Pig In Vivo. Eur. J. Pharmacol. (1989) 174(1):49-55	
BC	✓	IMAMURA, M. et al. Unmasking of Activated Histamine H3-Receptors in Myocardial Ischemia: Their Role as Regulators of Exocytotic Norepinephrine Release. J. Pharmacol. Exp. Ther. (1994) 271(3):1259-1266	
BC	✓	JONES, R.G. Studies on Imidazoles. II. The Synthesis of 5-Imidazolecarboxylates from Glycine and Substituted Glycine Esters. J. Am. Chem. Soc. (1949) 71:644-647	
BC	✓	JORDAAN, A.; ARNDT, R.R. The Synthesis of 1-Methyl-5-(α -indolyl)imidazole and 1-Methyl-2-ethylthiol-5-(α -indolyl)imidazole. Journal of Heterocyclic Chemistry 5(5), 723-5 (English) 1968	
BC	✓	KAPETANOVIC, I.M.; KUPFERBERG, H.J. Nafimidone, an Imidazole Anticonvulsant, and Its Metabolite as Potent Inhibitors of Microsomal Metabolism of Phenytoin and Carbamazepine. Drug Metab. Dispos. (1984) 12(5):580-584	
BC	✓	KORTE, A. et al. Characterization and Tissue Distribution of H3 Histamine Receptors in Guinea Pigs by N alpha-Methylhistamine. Biochem. Biophys. Res. Commun. (May 1990) 168(3):979-986	
BC	✓	KRAUSE, M. et al. Medicinal Chemistry of Histamine H3 Receptor Agonists; In The Histamine H3 Receptor - A Target for New Drugs Leurs, R.; Timmerman, H. (Eds.) Elsevier (1998) 175-198	
BC	✓	LAVRIJSEN, K. et al. Induction Potential of Antifungals Containing an Imidazole or Triazole Moiety. Biochem. Pharmacol. (1988) 35(11):1867-1878	
BC	✓	LEURS, R. et al. The Medicinal Chemistry and Therapeutic Potentials of Ligands of the Histamine H3 Receptor. Prog. Drug Res. (1995) 45:107-165	
BC	✓	LEURS, R. et al. "Therapeutic potential of histamine H3 receptor agonists and antagonists" Trends in Pharmacological sciences, Elsevier Trends Journal, Cambridge, BG, vol. 19, no. 5, 1 May 1998; Pages 177-184, XP004121095	
BC	✓	LIN, J.-S. et al. Involvement of Histaminergic Neurons in Arousal Mechanisms Demonstrated with H3-Receptor Ligands in the Cat. Brain Res. (1990) 523:325-330	
BC	✓	LINNEY, I.D. et al. Design, Synthesis, and Structure-Activity Relationships of Novel Non-Imidazole Histamine H3 Receptor Antagonists. J. Med. Chem. (2000) 43(12):2362-2370	
BC	✓	LOVENBERG, T.W. et al. Cloning and Functional Expression of the Human Histamine H3 Receptor. Mol. Pharmacol. (1999) 55:1101-1107	
BC	✓	LOVENBERG, T.W. et al. Cloning of Rat Histamine H3 Receptor Reveals Distinct Species Pharmacological Profiles. J. Pharmacol. Exp. Ther. (2000) 293(3):771-778	
BC	✓	MACHIDORI, H. et al. Zucker Obese Rats: Defect in Brain Histamine Control of Feeding. Brain Res. (1992) 590:180-188	
BC	✓	MCLEOD, R.L. et al. Antimigraine and Sedative Activity of SCH 50971: A Novel Orally-Active Histamine H3 Receptor Agonist. Soc. Neurosci. Abstr. (1998) 22:2010	
BC	✓	MEIER, G. et al. Piperidino-Hydrocarbon Compounds as Novel Non-Imidazole Histamine H3-Receptor Antagonists. Bioorg. Med. Chem. (2002) 10:2535-2542	
BC	✓	MONTI, J.M. et al. Effects of Selective Activation or Blockade of the Histamine H3 Receptor on Sleep and Wakefulness. Eur. J. Pharmacol. (1991) 205(3):283-287	
BC	✓	MORISSET, S. et al. High Constitutive Activity of Native H3 Receptors Regulates Histamine Neurons in Brain. Nature (Dec. 2000) 408:860-864	
BC	✓	ODA, T. et al. Molecular Cloning and Characterization of a Novel Type of Histamine Receptor Preferentially Expressed in Leukocytes. J. Biol. Chem. (2000) 275(47):36781-36786	
BC	✓	PANULA, P. et al. Significant Changes in the Human Brain Histaminergic System in Alzheimer's Disease. Soc. Neurosci. Abstr. (1995) 21:1977	



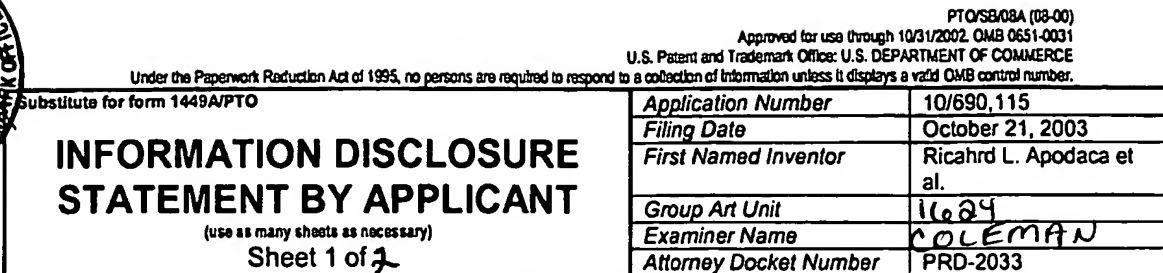
BC	✓	PHILLIPS, M.E. Positron Emission Tomography Provides Molecular Imaging of Biological Processes. Proc. Natl. Acad. Sci. (2000) 97(18):9226-9233
BC	✓	PHILLIPS, J.G.; ALI, S.M. Medicinal Chemistry of Histamine H3 Receptor Antagonists; In The Histamine H3 Receptor - A Target for New Drugs Leurs, R.; Timmerman, H. (Eds.) Elsevier (1998) 197-222
BC	✓	PHILLIPS, J.G. et al. Chapter 4, Recent Advances in Histamine H ₃ Receptor Agents. Ann. Reports in Med. Chem., 31, 1998, pages 31-40
BC	✓	ROULEAU, A. et al. Bioavailability, Antinociceptive and Antiinflammatory Properties of BP 2-94, a Histamine H3 Receptor Agonist Prodrug. J. Pharmacol. Exp. Ther. (1997) 281(3):1085-1094
BC	✓	SABBATINI, RENATO, M.E., The Cyclotron and PET. In Brain & Mind an electronic magazine about Neuroscience [online], March, 1997. Retrived from the internet, <http://www.epub.org.br/cm/n01/pe/pet/cyclo.htm>
BC	✓	SCHLICKER, E.; MARR, I. The Moderate Affinity of Clozapine at H3 Receptors Is Not Shared by Its Two Major Metabolites and by Structurally Related and Unrelated Atypical Neuroleptics. Naunyn-Schmiedeberg's Arch. Pharmacol. (1998) 353:290-294
BC	✓	SHEETS, J.J.; MASON, J.I. Ketoconazole: a Potent Inhibitor of Cytochrome P-450-Dependent Drug Metabolism in Rat Liver. Drug Metab. Dispos. (1984) 12(5):603-608
BC	✓	STARK, H. et al. Developments of Histamine H3-Receptor Antagonists. Drugs Future (1996) 21(5):507-520
BC	✓	TOZER, M.J., et al.: "From Histamine to imidazolylalkyl-sulfonamides: the design of a novel series of histamine H3 receptor antagonists"; Bioorganic & Medicinal Chemistry Letters, OXFORD, GB, vol. 9, no. 13, 5 July 1999, Pages:1825-1830, XP004168848
BC	✓	TOZER, M.J.; KALINDJIAN, S.B. Histamine H3 Receptor Antagonists. Exp. Opin. Ther. Patents (2000) 10(7):1045-1055
BC	✓	WALCZYNSKI, K. et al. Non-Imidazole Histamine H3 Ligands, Part 2: New 2-Substituted Benzothiazoles as Histamine H3 Antagonists. Arch. Pharm. Pharm. Med. Chem. (Weinheim, Ger.) (1999) 332:389-398
BC	✓	WALCZYNSKI, K. et al. Non-Imidazole Histamine H3 Ligands. Part 1. Synthesis of 2-(1-Piperazinyl)- and 2-(Hexahydro-1H-1,4-diazepin-1-yl)benzothiazole Derivatives as H3-Antagonists with H1 Blocking Activities. Farmaco (1999) 54:684-694
BC	✓	WEST, R.E. et al. Identification of Two H3-Histamine Receptor Subtypes. Mol. Pharmacol. (1990) 38(5):610-613
BC	✓	WEST, R.E., Jr. et al. The Profiles of Human and Primate [3H]N alpha-methylhistamine Binding Differ from That of Rodents. Eur. J. Pharmacol. (1999) 377:233-239
BC	✓	YOKOYAMA, H. et al. Effect of Thioperamide, a Histamine H3 Receptor Antagonist, on Electrically Induced Convulsions in Mice. Eur. J. Pharmacol. (1993) 234:129-133
BC	✓	ANJANEYULU, B. et al. Synthesis of 14C-Labelled 1-Methanesulphonyl-3-(1-methyl-5-nitro-1H-imidazol-2-yl)-2-imidazolidinone, (Go 10213). J. Labelled Compd. Radiopharm. (1983) 20(8):951-961
BC	✓	ITEMURA, R. et al. Synthesis of Benzimidazole Derivatives as Potential H1-Antihistaminic Agents. J. Heterocycl. Chem. (1987) 24:31-37
BC	✓	IWATA, R. et al. Synthesis of 3-[1H-imidazol-4-yl]propyl 4-[18F]fluorobenzyl Ether ([18F]Fluoroproxyfan): A Potential Radioligand for Imaging Histamine H3 Receptors. J. Labelled Compd. Radiopharm. (2000) 43:873-882
BC	✓	JAROSINSKI, M.A.; ANDERSON, W.K. Preparation of Noncondensed 2-Substituted 1-Methylimidazoles via Ipsso Substitution Reaction on 2-Sulfinyl or 2-Sulfonyl Derivatives of 4,5-Disubstituted 1-Methylimidazoles. J. Org. Chem. (1991) 56(12):4058-4062
BC	✓	OHTA, S. et al. Synthesis and Application of Imidazole Derivatives. Introduction of Carbogenic Substituents into the 5-Position of 1-Methyl-1H-imidazole. Chem. Pharm. Bull. (1992) 40(10):2681-2685
BC	✓	PHILLIPS, B.T. et al. Preparation of 5-Substituted 2-Mercapto-1-methylimidazoles. Direct Metalation of 2-Mercapto-1-methylimidazole. Synthesis (1990):781-783
BC	✓	SCHNETTLER, R.A. et al. 4-Aroyl-1,3-dihydro-2H-imidazol-2-ones, a New Class of Cardiotonic Agents. J. Med. Chem. (1982) 25:1477-1481
BC	✓	SHAPIRO, G.; MARZI, M. Synthesis of 2,5-Dilithio-1-methylimidazole. Tetrahedron Lett. (1993) 34(21):3401-3404
BC	✓	ERDELYI, M.; GOGOLL, A. Rapid Homogeneous-Phase Sonogashira Coupling Reactions Using Controlled Microwave Heating. J. Org. Chem. (2001) 66(12):4165-4169
BC	✓	APODACA, R. et al. A New Class of Diamine-based Histamine H3 Receptor Antagonists: 4-(Aminoalkoxy)benzylamines. J. Med. Chem. (2003) 46(18):3938-3944
BC	✓	STARK, H. Recent Advances in Histamine H3/H4 Receptor Ligands. Expert Opin. Ther. Patents (2003) 13(6):851-865
BC	✓	Phenylalkynes to Treat Histamine-Mediated Conditions. Expert Opin. Ther. Patents (2003) 13(11):1759-1762

Examiner Signature	Brenda Coleman	Date Considered	May 30, 2006
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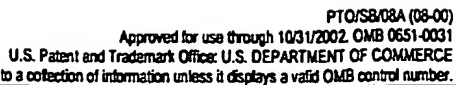
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